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NEWS	1		Web Page URLs for STN Seminar Schedule - N. America
NEWS	2		"Ask CAS" for self-help around the clock
NEWS	3	SEP 01	New pricing for the Save Answers for SciFinder Wizard within STN Express with Discover!
NEWS	4	OCT 28	KOREAPAT now available on STN
NEWS	5	NOV 30	PHAR reloaded with additional data
NEWS	6	DEC 01	LISA now available on STN
NEWS	7	DEC 09	12 databases to be removed from STN on December 31, 2004
NEWS	8	DEC 15	MEDLINE update schedule for December 2004
NEWS	9	DEC 17	ELCOM reloaded; updating to resume; current-awareness alerts (SDIs) affected
NEWS	10	DEC 17	COMPUAB reloaded; updating to resume; current-awareness alerts (SDIs) affected
NEWS	11	DEC 17	SOLIDSTATE reloaded; updating to resume; current-awareness alerts (SDIs) affected
NEWS	12	DEC 17	CERAB reloaded; updating to resume; current-awareness alerts (SDIs) affected
NEWS	13	DEC 17	THREE NEW FIELDS ADDED TO IFIPAT/IFIUDB/IFICDB
NEWS	14	DEC 30	EPFULL: New patent full text database to be available on STN
NEWS	15	DEC 30	CAPLUS - PATENT COVERAGE EXPANDED
NEWS	16	JAN 03	No connect-hour charges in EPFULL during January and February 2005
NEWS	17	JAN 11	CA/CAPLUS - Expanded patent coverage to include Russia (Federal Institute of Industrial Property)

NEWS EXPRESS JANUARY 10 CURRENT WINDOWS VERSION IS V7.01a, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP), AND CURRENT DISCOVER FILE IS DATED 10 JANUARY 2005

NEWS HOURS	STN Operating Hours Plus Help Desk Availability
NEWS INTER	General Internet Information
NEWS LOGIN	Welcome Banner and News Items
NEWS PHONE	Direct Dial and Telecommunication Network Access to STN
NEWS WWW	CAS World Wide Web Site (general information)

Enter NEWS followed by the item number or name to see news on that specific topic.

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  * * * * * * * * * * * * * * STN  Columbus  * * * * * * * * * * * * * * *

FILE 'HOME' ENTERED AT 09:20:34 ON 13 JAN 2005

=> FIL STNGUIDE
COST IN U.S. DOLLARS                                    SINCE FILE      TOTAL
                                                           ENTRY    SESSION
FULL ESTIMATED COST                                         0.21       0.21

FILE 'STNGUIDE' ENTERED AT 09:20:47 ON 13 JAN 2005
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AND TECHNOLOGY CORPORATION, AND FACHINFORMATIONSZENTRUM KARLSRUHE

FILE CONTAINS CURRENT INFORMATION.
LAST RELOADED: Jan  7, 2005 (20050107/UP).

=> FIL HOME
COST IN U.S. DOLLARS                                    SINCE FILE      TOTAL
                                                           ENTRY    SESSION
FULL ESTIMATED COST                                         0.06       0.27

FILE 'HOME' ENTERED AT 09:20:51 ON 13 JAN 2005

=> file reg
COST IN U.S. DOLLARS                                    SINCE FILE      TOTAL
                                                           ENTRY    SESSION
FULL ESTIMATED COST                                         0.21       0.48

FILE 'REGISTRY' ENTERED AT 09:20:55 ON 13 JAN 2005
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Property values tagged with IC are from the ZIC/VINITI data file
provided by InfoChem.

STRUCTURE FILE UPDATES:   11 JAN 2005  HIGHEST RN 811782-89-5
DICTIONARY FILE UPDATES:  11 JAN 2005  HIGHEST RN 811782-89-5

TSCA INFORMATION NOW CURRENT THROUGH MAY 21, 2004

  Please note that search-term pricing does apply when
  conducting SmartSELECT searches.

Crossover limits have been increased.  See HELP CROSSOVER for details.

Experimental and calculated property data are now available. For more
information enter HELP PROP at an arrow prompt in the file or refer
to the file summary sheet on the web at:
http://www.cas.org/ONLINE/DBSS/registryss.html

=>
Uploading C:\STNEXP4\QUERIES\10824826.str
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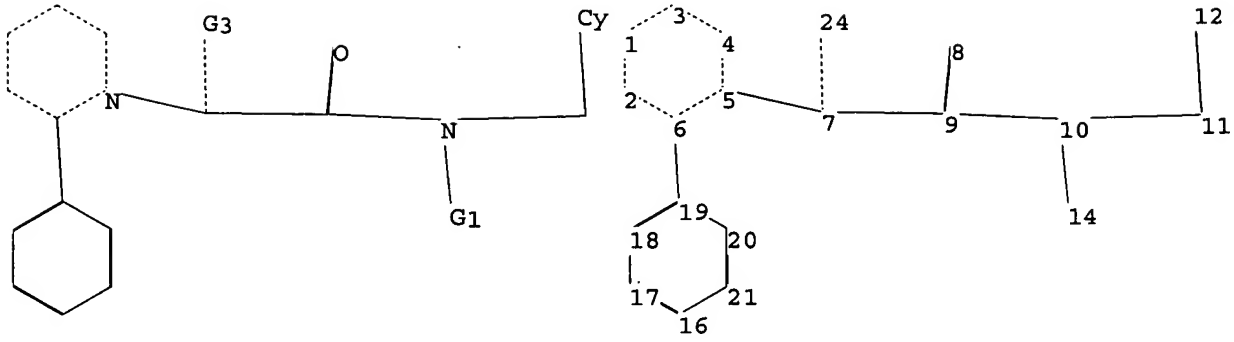


chain nodes :
7 8 9 10 11 12 14 24
ring nodes :
1 2 3 4 5 6 16 17 18 19 20 21
chain bonds :
5-7 6-19 7-24 7-9 8-9 9-10 10-11 10-14 11-12
ring bonds :
1-3 1-2 2-6 3-4 4-5 5-6 16-17 16-21 17-18 18-19 19-20 20-21
exact/norm bonds :
1-3 1-2 2-6 3-4 4-5 5-6 5-7 7-24 8-9 9-10 10-11 10-14 11-12
exact bonds :
6-19 7-9
normalized bonds :
16-17 16-21 17-18 18-19 19-20 20-21
isolated ring systems :
containing 1 : 16 :

G1:Cy,Ak

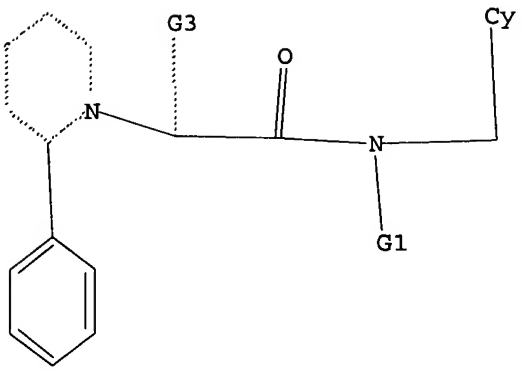
G2:Cy,Ak

G3:X,Ak,H

Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom
11:Atom 12:Atom 14:CLASS 16:Atom 17:Atom 18:Atom 19:Atom 20:Atom 21:Atom
24:CLASS
Generic attributes :
12:
Saturation : Unsaturated

L1 STRUCTURE UPLOADED

=> dis l1
L1 HAS NO ANSWERS
L1 STR



G1 Cy,Ak

G2 Cy,Ak

G3 X,Ak,H

Structure attributes must be viewed using STN Express query preparation.

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=> s l1 sam
SAMPLE SEARCH INITIATED 09:21:15 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED -    521 TO ITERATE

100.0% PROCESSED     521 ITERATIONS                                1 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS:  ONLINE  **COMPLETE**
                        BATCH   **COMPLETE**
PROJECTED ITERATIONS:          9051 TO    11789
PROJECTED ANSWERS:                1 TO       80

L2             1 SEA SSS SAM L1

=> s l1 full
FULL SEARCH INITIATED 09:21:20 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED -  10285 TO ITERATE

100.0% PROCESSED   10285 ITERATIONS                                9 ANSWERS
SEARCH TIME: 00.00.02

L3             9 SEA SSS FUL L1

=> file hcaplus
COST IN U.S. DOLLARS                                SINCE FILE      TOTAL
                                                       ENTRY    SESSION
FULL ESTIMATED COST                                   161.33     161.81

FILE 'HCAPLUS' ENTERED AT 09:21:30 ON 13 JAN 2005
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FILE COVERS 1907 - 13 Jan 2005 VOL 142 ISS 3
FILE LAST UPDATED: 12 Jan 2005 (20050112/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s l3
L4 2 L3

=> dis l4 1-2 bib abs hitstr

L4 ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2005 ACS on STN
AN 2003:796667 HCAPLUS
DN 139:307693
TI Preparation of substituted tetrahydroisoquinolines as C5a receptor modulators
IN Mitchell, Scott; Ohliger, Robert; Zhang, Luyan; Zhao, He; Currie, Kevin; Lee, Kyungae
PA Neurogen Corporation, USA
SO PCT Int. Appl., 104 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003082828	A1	20031009	WO 2003-US9046	20030325
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	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	EP 1487798	A1	20041222	EP 2003-714371	20030325
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
	US 2004006069	A1	20040108	US 2003-401135	20030327
	US 6777422	B2	20040817		
	US 2004204446	A1	20041014	US 2004-824826	20040415
PRAI	US 2002-368199P	P	20020328		
	WO 2003-US9046	W	20030325		
	US 2003-401135	A1	20030327		

OS MARPAT 139:307693
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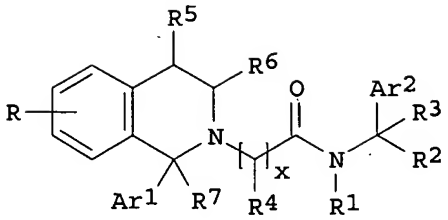

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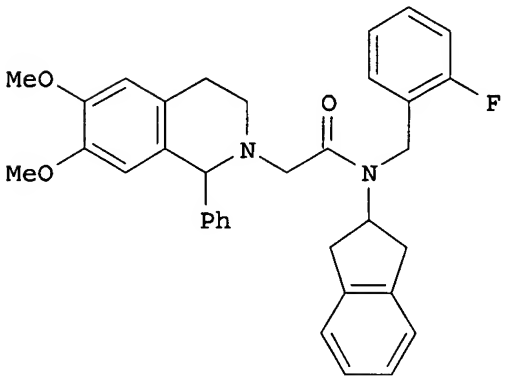


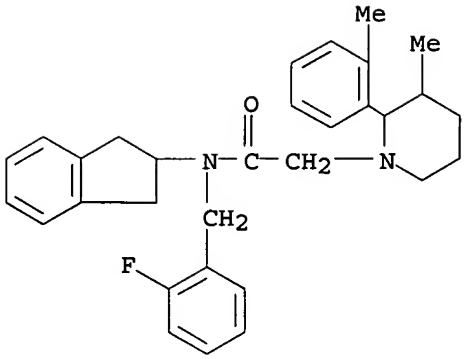
II

AB The title compds. [I; x = 1-3; R = halo, OH, alkoxy, etc.; R1 = alkyl, alkenyl, cycloalkyl, etc.; R2-R4 = H, halo, alkyl, alkoxy; R5, R6 = H, halo, OH, etc.; R7 = H, alkyl, alkenyl, etc.; Ar1 = (un)substituted Ph, naphthyl, biphenyl, etc.; Ar2 = (un)unsubstituted aryl, heteroaryl] which are ligands that may be used to modulate C5a receptor activity in vivo or in vitro, and are particularly useful in the treatment of conditions associated with pathol. C5a receptor activation in humans, domesticated companion animals and livestock animals, were prepared Thus, reacting 6,7-dimethoxy-1-phenyl-1,2,3,4-tetrahydroisoquinoline.HCl with N-(1-fluorobenzyl)-N-(indan-2-yl)-2-bromoacetamide in the presence of K2CO3 in MeCN afforded II. Preferred compds. I exhibit IC50 values of less than 1 μM in the assay for C5a receptor mediated chemotaxis. Pharmaceutical compns. and methods for using them to treat disorders associated with pathol. C5a receptor activation are provided, as are methods for using such ligands for receptor localization studies.

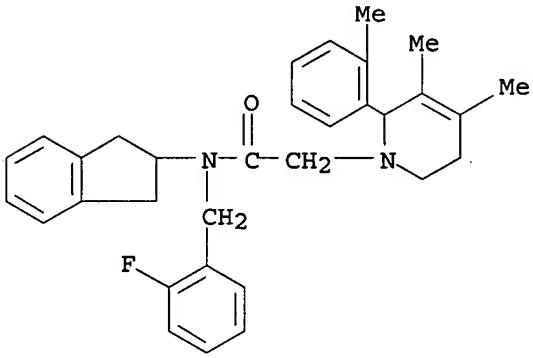
IT **610296-34-9P 610296-35-0P 610298-33-4P**
610298-35-6P 610298-39-0P 610298-43-6P
610298-45-8P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of new aryl imidazoles and related compds. as C5a receptor modulators)

RN 610296-34-9 HCAPLUS

CN 1-Piperidineacetamide, N-(2,3-dihydro-1H-inden-2-yl)-N-[(2-fluorophenyl)methyl]-3-methyl-2-(2-methylphenyl)- (9CI) (CA INDEX NAME)

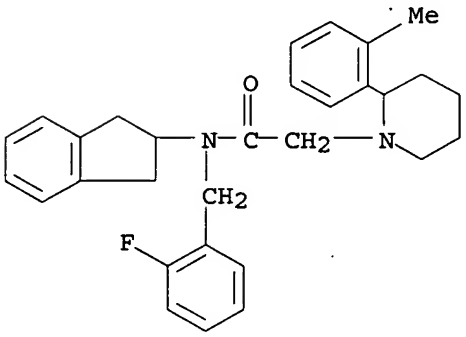


RN 610296-35-0 HCAPLUS
CN 1(2H)-Pyridineacetamide, N-(2,3-dihydro-1H-inden-2-yl)-N-[(2-fluorophenyl)methyl]-5,6-dihydro-3,4-dimethyl-2-(2-methylphenyl)-, monohydrochloride (9CI) (CA INDEX NAME)

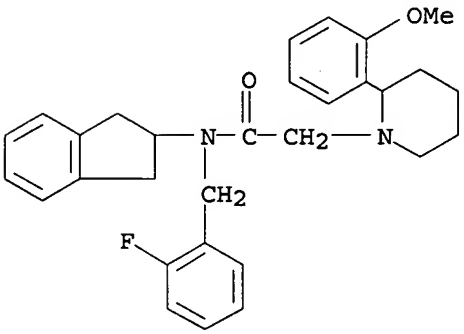


● HCl

RN 610298-33-4 HCAPLUS
CN 1-Piperidineacetamide, N-(2,3-dihydro-1H-inden-2-yl)-N-[(2-fluorophenyl)methyl]-2-(2-methylphenyl)- (9CI) (CA INDEX NAME)

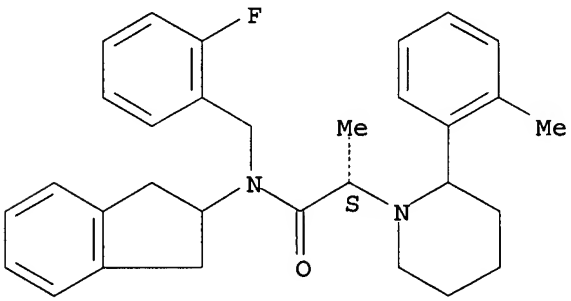


RN 610298-35-6 HCAPLUS
CN 1-Piperidineacetamide, N-(2,3-dihydro-1H-inden-2-yl)-N-[(2-fluorophenyl)methyl]-2-(2-methoxyphenyl)- (9CI) (CA INDEX NAME)

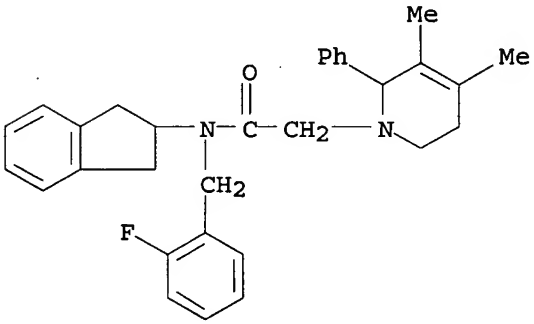


RN 610298-39-0 HCAPLUS
CN 1-Piperidineacetamide, N-(2,3-dihydro-1H-inden-2-yl)-N-[(2-fluorophenyl)methyl]-α-methyl-2-(2-methylphenyl)-, (αS)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

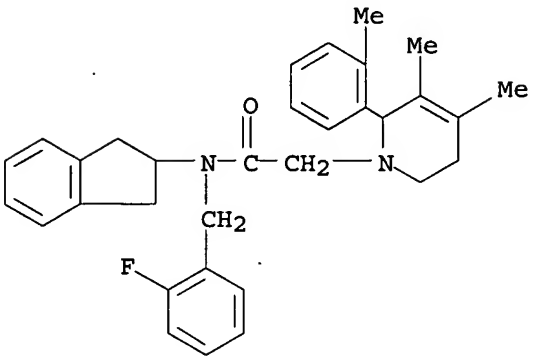


RN 610298-43-6 HCAPLUS
CN 1(2H)-Pyridineacetamide, N-(2,3-dihydro-1H-inden-2-yl)-N-[(2-fluorophenyl)methyl]-5,6-dihydro-3,4-dimethyl-2-phenyl- (9CI) (CA INDEX NAME)



RN 610298-45-8 HCAPLUS

CN 1(2H)-Pyridineacetamide, N-(2,3-dihydro-1H-inden-2-yl)-N-[(2-fluorophenyl)methyl]-5,6-dihydro-3,4-dimethyl-2-(2-methylphenyl)- (9CI) (CA INDEX NAME)



RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2005 ACS on STN
AN 1994:435340 HCAPLUS
DN 121:35340
TI Preparation of azacyclic compounds as tachykinin antagonists.
IN Seward, Eileen Mary; Swain, Christopher John
PA Merck Sharp and Dohme Ltd., UK
SO PCT Int. Appl., 54 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9402461	A1	19940203	WO 1993-GB1525	19930720
	W: AU, CA, JP, US				
	RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
	EP 652866	A1	19950517	EP 1993-917877	19930720
	EP 652866	B1	19981125		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, NL, PT, SE				
	JP 07508993	T2	19951005	JP 1993-503945	19930720
	AU 675447	B2	19970206	AU 1993-47139	19930720
	AU 9347139	A1	19940214		
	AT 173725	E	19981215	AT 1993-917877	19930720
	ES 2124318	T3	19990201	ES 1993-917877	19930720
	US 5561130	A	19961001	US 1995-379622	19950124
	US 5496833	A	19960305	US 1995-387684	19950213
PRAI	GB 1992-16065	A	19920728		
	GB 1992-16304	A	19920731		
	GB 1992-24918	A	19921127		
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	US 1993-46538	A3	19930413		
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OS MARPAT 121:35340
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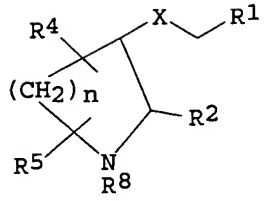


I

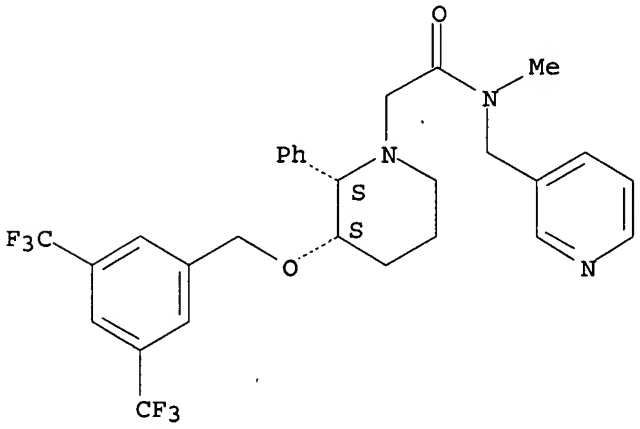
AB Title compds. [I; n = 1-3; X = O, S; R1 = (substituted) Ph; R2 = (substituted) aryl, heteroaryl; R4, R5 = H, halo, C1-6 alkyl, oxo, CH2ORa, CO2Ra or CONRaRb; R8 = C(COORa)2, C(CONRaRb)2 or C1-6 alkyl substituted by C(:NRa)NRbNRcCO2Rd, CONHNRaRb, C(S)NRaRb, etc.; Ra, Rb, Rc, Rd = H, C1-6 alkyl, Ph, trifluoromethyl], were prepared with claimed uses of treating/preventing pain, inflammation, migraine, or arthritis. Thus, (2R*,3R*)-3-[(3,5-bis(trifluoromethyl)phenyl)methoxy]-1-(carboxymethyl)-2-phenylpiperidine (preparation given) was stirred with 1-hydroxybenzotriazole, 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride, Et3N, and propargylamine in THF overnight at room temperature to give (2R*,3R*)-3-[(3,5-bis(trifluoromethyl)phenyl)methoxy]-2-phenyl-1-[N-(prop-2-ynyl)carboxamidomethyl]piperidine. I inhibited substance P binding to human NK1R with IC50 ≤100 nM. Generic I formulations are given.

IT **155847-34-0P 155847-46-4P**
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of, as tachykinin antagonist)

RN 155847-34-0 HCAPLUS

CN 1-Piperidineacetamide, 3-[[3,5-bis(trifluoromethyl)phenyl]methoxy]-N-methyl-2-phenyl-N-(3-pyridinylmethyl)-, (2S-cis)- (9CI) (CA INDEX NAME)

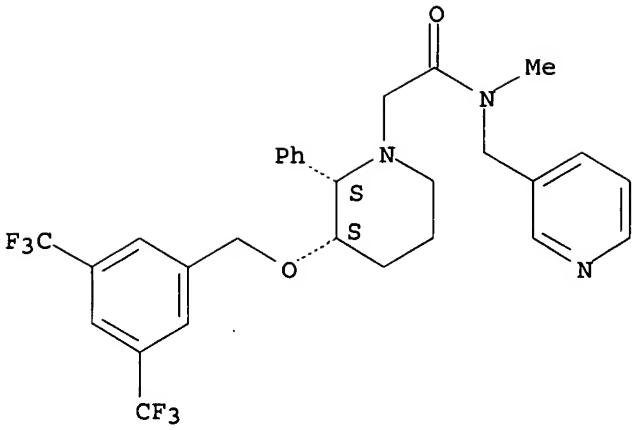
Absolute stereochemistry.



RN 155847-46-4 HCAPLUS

CN 1-Piperidineacetamide, 3-[[3,5-bis(trifluoromethyl)phenyl]methoxy]-N-methyl-2-phenyl-N-(3-pyridinylmethyl)-, monohydrobromide, (2S-cis)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



● HBr

=> log y

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	12.33	174.14

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-1.46	-1.46

STN INTERNATIONAL LOGOFF AT 09:21:50 ON 13 JAN 2005